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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
10/709,413	05/04/2004	Yu-Jie Zhao	46863	3412
31561	7590 09/30/2005		EXAMINER	
JIANQ CHYUN INTELLECTUAL PROPERTY OFFICE			YU, MELANIE J	
7 FLOOR-1, 1 ROOSEVELT	NO. 100 ROAD, SECTION 2		ART UNIT	PAPER NUMBER
TAIPEI, 100 TAIWAN		1641		
			DATE MAILED: 09/30/2005	

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)				
Office Action Summary		10/709,413	ZHAO, YU-JIE	,			
		Examiner	Art Unit				
		Melanie Yu	1641				
Period fo	The MAILING DATE of this communication apports and the second	pears on the cover sheet with the	correspondence address				
WHIC - Exte after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLEMENTER IS LONGER, FROM THE MAILING Densions of time may be available under the provisions of 37 CFR 1.1 SIX (6) MONTHS from the mailing date of this communication. Operiod for reply is specified above, the maximum statutory period are to reply within the set or extended period for reply will, by statuted reply received by the Office later than three months after the mailing patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 136(a). In no event, however, may a reply be will apply and will expire SIX (6) MONTHS from a cause the application to become ABANDON	ON. timely filed om the mailing date of this communication NED (35 U.S.C. § 133).				
Status							
1)	Responsive to communication(s) filed on 14 J	ulv 2005					
2a)∏	<u> </u>	s action is non-final.					
, <u> </u>	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is						
,	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Dispositi	ion of Claims						
•	4) Claim(s) 1-9 is/are pending in the application.						
	4a) Of the above claim(s) is/are withdra Claim(s) is/are allowed.	wil from consideration.					
	Claim(s) is/are allowed. Claim(s) <u>1-9</u> is/are rejected.						
<u> </u>	Claim(s) is/are objected to.						
	Claim(s) are subject to restriction and/o	or election requirement					
		or cicotion requirement.					
Applicati	on Papers						
<u> </u>	The specification is objected to by the Examine						
10)⊠	The drawing(s) filed on <u>04 May 2005</u> is/are: a)	•	•				
	Applicant may not request that any objection to the	•	` '				
445	Replacement drawing sheet(s) including the correct		·	d).			
11)[The oath or declaration is objected to by the Ex	xaminer. Note the attached Office	e Action or form PTO-152.				
Priority ι	ınder 35 U.S.C. § 119						
a)	Acknowledgment is made of a claim for foreign All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureasee the attached detailed Office action for a list	ts have been received. Is have been received in Application of the contraction of the co	ation No ved in this National Stage				
2) Notic 3) Inform	t(s) e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449 or PTO/SB/08) r No(s)/Mail Date	4) Interview Summa Paper No(s)/Mail 5) Notice of Informal 6) Other:					

DETAILED ACTION

Applicant's amendment filed 14 July 2005 has been entered. Claims 1, 4 and 5 are 1. currently amended. Claims 1-9 are currently pending in this application. Claims 10-20 are cancelled.

Withdrawn Rejections

Previous rejection of claims 1-9 under 35 USC 112, second paragraph, 35 USC 102(b) and 35 USC 103(a) have been withdrawn in light of applicant's arguments and amendments.

Claim Rejections - 35 USC § 102

2. Claims 1 and 2 are rejected under 35 U.S.C. 102(e) as being anticipated by Blackburn (US 2003/0190608).

Blackburn teaches a method of fabricating a cell detection chip, comprising: selecting a plurality of probe molecules, wherein an affinity exists between each of the probe molecules and one of corresponding antigens on a cell membrane (different capture probes are specific for analyte, par. 150; analyte are antigens on cell membranes, par. 103); modifying the plurality of probe molecules to facility an immobilization of the probe molecules onto a matrix (par. 162); and spotting the probe molecules respectively onto respective positions of the matrix (par. 152-153).

Claim Rejections - 35 USC § 103

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

3. Claims 1, 2, 4-6 and 9 are rejected under 35 U.S.C. 102(b) as being anticipated by Okamoto et al. (US 2003/0059817) in view of Kapur et al. (US 6,548,263).

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With respect to claims 1, 2 and 6, Okamoto et al. teach a method of fabricating a cell detection chip, comprising: designing a plurality of probe molecules, wherein an affinity exists between each of the probe molecules and one of corresponding specific molecules (par. 0056); synthesizing a plurality of probe molecules (par. 0046, 0056); spotting the probe molecules respectively on a matrix (par. 0056); and incubating the matrix to keep the matrix under a wet environment (support stood in a humid chamber for 30 minutes, par. 0056). Okamoto et al. fail to teach specific molecules being on a cell membrane.

Blackburn teaches a spotted array comprising probe molecules (col. 13, lines 50-67; col. 14, lines 53-55) wherein the corresponding specific molecule (analyte) is an antigen on a cell membrane (col. 15, lines 62-67), in order to provide a high throughput specific cell-type binding microarray.

Therefore it would have been obvious to one having ordinary skill in the art at the time the invention was made to include in the method of Okamoto et al., specific molecules on the surface of cell membranes as taught by Kapur et al., in order to provide high biological content screening for drug candidates by analysis of drug-cell interactions when a small number of cells and large volumes of compounds required for testing.

Regarding claims 4 and 5, Okamoto et al. teach designing probe molecules comprising a plurality of location indication probes (par. 0118) and the step of synthesizing the probe molecules, further comprising the step of dissolving probe molecules in a solvent to form a solution of the probe molecules (probe molecules are mixed in a solution, par. 0056).

With respect to claim 9, Okamoto et al. teach a spot diameter between 20 and 100 μ m (par. 0033), which encompasses the recited range of a spot radius between 50 and 500 μ m.

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4. Claim 3 is rejected under 35 U.S.C. 103(a) as being unpatentable over Okamoto et al. (US 2003/0059817) in view of Kapur et al. (US 6,548,263), as applied to claim 1, and further in view of Chen et al. (US 6,594,432).

Okamoto et al. in view of Kapur et al., as applied to claim 1, teach a method of fabricating a cell detection chip, but fail to teach the step of designing probe molecules further comprising designing a plurality of quality control probes.

Chen et al. teach using a plurality of quality control probes (col. 7, lines 10-22), in order to inspect microarrays after their formation.

Therefore it would have been obvious to on having ordinary skill in the art at the time the invention was made to include in the designing step of the method of Okamoto et al. in view of Kapur et al., designing a plurality of quality control probes as taught by Chen et al., in order to determine if probes have been deposited.

5. Claims 7 and 8 are rejected under 35 U.S.C. 103(a) as being unpatentable over Okamoto et al. (US 2003/0059817) in view of Kapur et al. (US 6,548,263) further in view of Oprandy (US 5,200,312).

Okamoto et al. in view of Kapur et al., as applied to claims 1 and 6, teach a method of fabricating a cell detection chip and a step of cleaning after incubation (par. 0116), but fail to teach a step of drying after an incubation step and before cleaning.

Oprandy teaches a step of drying (col. 4, lines 11-19), in order to store an antibody bound membrane for later use.

Therefore it would have been obvious to one having ordinary skill in the art at the time the invention was made to include in the method of fabricating a chip after the step of incubation Art Unit: 1641

and the step of cleaning of Okamoto et al. in view of Kapur et al., a step of drying as taught by Oprandy, in order to ensure the probe has completely bound to the matrix.

With respect to claim 8, Okamoto et al. teach after the step of cleaning, steps of: blocking portions of a surface of the matrix not spotted with probes, wherein a blocking solution is used (immersed in bovine serum albumin to proceed blocking reaction, par. 0116); and further cleaning the matrix (matrix is washed after hybridization reaction, par. 0118).

Response to Arguments

Applicant's arguments, see pages 5-7, filed 14 July 2005, with respect to the rejection(s) 6. of claim(s) 1-9 under 35 USC 112, second paragraph and 35 USC 102(b) have been fully considered and are persuasive. Therefore, the rejection has been withdrawn. However, upon further consideration, a new ground(s) of rejection is made in view of a specific molecule present on a cell membrane.

Conclusion

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Melanie Yu whose telephone number is (571) 272-2933. The examiner can normally be reached on M-F 8:30-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on (571) 272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Melanie Yu

Patent Examiner

Melanie

Art Unit 1641

LONG V. LE

SUPERVISORY PATENT EXAMINER

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